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Amendments to the Claims:

Please amend claims 1, 5 and 8. This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

- 1. (Currently Amended) A computer-based method for identifying conserved peptide motifs useful as drug targets for use in a host organism, wherein the said method comprises the steps of:
- i) providing electronic data representing peptide libraries from the protein sequences of selected organisms,
- ii) from the data of step (i), generating computationally overlapping peptide sequences from selected organisms of length 'N',-and
- (ii) sorting computationally the peptide sequences of length 'N' according to amino acid sequence,
- (iii) matching computationally the sorted peptide sequences of length 'N' of the selected organisms to produce matched common peptide sequences,
- (iv) locating computationally the matched common peptide sequences in the protein sequences of step i) and subsequently labeling the matched common peptide sequences with their origin and location,
- (v) joining computationally overlapping common peptide sequences to obtain extended conserved peptide sequences,
- (vi) annotating secondary structure of extended conserved peptide sequences based on a crystal structure database, and
- vii) comparing known proteins of a pathogenic organism with those of non-pathogenic organisms using the aforementioned steps (i) to (v), to select at least one conserved peptide

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sequence not commonly conserved in both the pathogenic organism and in non-pathogenic organisms, to obtain a conserved peptide sequence,

viii) validating computationaly the conserved peptide sequences obtained in step (vii) as a potential drug target sequences by searching for the conserved peptide sequence in a host organism identifying conserved sequences not present in the host organism.

- 2. (Previously Presented) The method of claim 1 wherein 'N' is at least 4.
- 3. (Previously Presented) The method of claim 1 wherein the selected organisms include at least one of: Mycoplasma pneumoniae, Helicobacter pylori, Hemophillus influenzae, Mycobacterium tuberculosis, Mycoplasma genitalium, Bacillus subtillis, and Escherichia coli.
- 4. (Previously Presented) A method as claimed in claim 1 where conserved peptide motifs as modified comprising sequences include one or more of:

| | • • • | | |
|-----|----------------------------|-------|------------------------------|
| 1. | AAQSIGEPGTQLT (SEQ ID NO:1 |) 35. | KMSKSKGN (SEQ ID NO:35) |
| 2. | AGDGTTTAT (SEQ ID NO:2) | 36. | KMSKSLGN (SEQ ID NO:36) |
| 3. | AGRHGNKG (SEQ ID NO:3) | 37. | KNMITGAAQMDGAILVV (SEQ |
| | | | ID NO:37) |
| 4. | AHIDAGKTTT (SEQ ID NO:4) | 38. | KPNSALRK (SEQ ID NO:38) |
| 5. | CPIETPEG (SEQ ID NO:5) | 39. | LFGGAGVGKTV (SEQ ID NO:39) |
| 6. | DEPSIGLH (SEQ ID NO:6) | 40. | LGPSGCGK (SEQ ID NO:40) |
| 7. | DEPTSALD (SEQ ID NO:7) | 41. | LHAGGKFD (SEQ ID NO:41) |
| 8. | DEPTTALDVT (SEQ ID NO:8) | 42. | LIDEARTPLIISG (SEQ ID NO:42) |
| 9. | DHAGIATQ (SEQ ID NO:9) | 43. | LLNRAPTLH (SEQ ID NO:43) |
| 10. | DHPHGGGEG (SEQ ID NO10) | 44. | LPDKAIDLIDE (SEQ ID NO:44) |
| 11. | DLGGGTFD (SEQ ID NO:11) | 45. | LPGKLADC (SEQ ID NO:45) |
| 12. | DVLDTWFSS (SEQ ID NO:12) | 46. | LSGGQQQR (SEQ ID NO:46) |
| 13. | ERERGITI (SEQ ID NO:13) | 47. | MGHVDHGKT (SEQ ID NO:47) |

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| 14. | ERGITITSAAT (SEQ ID NO:14) | 48. | NADFDGDQMAVH (SEQ ID |
|-----|-----------------------------|------|-----------------------------|
| | | | NO:48) |
| 15. | ESRRIDNQLRGR (SEQ ID NO:15) | 49. | NGAGKSTL (SEQ ID NO:49) |
| 16. | FSGGQRQR (SEQ ID NO:16) | 50. | NLLGKRVD (SEQ ID NO:50) |
| 17. | GEPGVGKTA (SEQ ID NO:17) | 51. | NTDAEGRL (SEQ ID NO:51) |
| 18. | GFDYLRDN (SEQ ID NO:18) | 52. | PSAVGYQPTLA (SEQ ID NO:52) |
| 19. | GHNLQEHS (SEQ ID NO:19) | 53. | QRVALARA (SEQ ID NO:53) |
| 20. | GIDLGTTNS (SEQ ID NO:20) | 54. | QRYKGLGEM (SEQ ID NO:54) |
| 21. | GINLLREGLD (SEQ ID NO:21) | 55. | RDGLKPVHRR (SEQ ID NO:55) |
| 22. | GIVGLPNVGKS (SEQ ID NO:22) | 56. | SALDVSIQA (SEQ ID NO:56) |
| 23. | GKSSLLNA (SEQ ID NO:23) | 57. | SGGLHGVG (SEQ ID NO:57) |
| 24. | GLTGRKIIVDTYG(SEQ ID NO:24 |)58. | SGSGKSSL (SEQ ID NO:58) |
| 25. | GPPGTGKTLLA (SEQ ID NO:25) | 59. | SGSGKSTL (SEQ ID NO:59) |
| 26. | GPPGVGKT (SEQ ID NO:26) | 60. | SVFAGVGERTREGND (SEQ ID |
| | | | NO:60) |
| 27. | GSGKTTLL (SEQ ID NO:27) | 61. | TGRTHQIRVH (SEQ ID NO:61) |
| 28. | GTRIFGPV (SEQ ID NO:28) | 62. | TGVSGSGKS (SEQ ID NO:62) |
| 29. | IDTPGHVDFT (SEQ ID NO:29) | 63. | TLSGGEAQRI (SEQ ID NO:63) |
| 30. | ILAHIDHGKSTL (SEQ ID NO:30) | 64. | TNKYAEGYP (SEQ ID NO:64) |
| 31. | INGFGRIGR (SEQ ID NO:31) | 65. | TPRSNPATY (SEQ ID NO:65) |
| 32. | IREGGRTVG (SEQ ID NO:32) | 66. | VEGDSAGG (SEQ ID NO:66) and |
| 33. | IVGESGSGKS (SEQ ID NO:33) | 67. | VRKRPGMYIG (SEQ ID NO:67). |
| 34. | KFSTYATWWI (SEQ ID NO:34) | | |
| | | | |

(Currently Amended) A method as claimed in claim 1 comprising increasing the 5. number of [invariant] conserved peptide sequences by increasing the relatedness among the organisms being compared.

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6. (Previously Presented) A method as claimed in any one of claims 1-4 wherein the invariant sequences belong to at least one of the following proteins:

- I DNA DIRECTED RNA POLYMERASE BETA CHAIN
- II EXCINUCLEASE ABC SUBUNIT A
- III EXCINUCLEASE ABC SUBUNIT B
- IV DNA GYRASE SUBUNIT B
- V ATP SYNTHASE BETA CHAIN
- VI S-ADENOSYLMETHIONINE SYNTHETASE
- VII GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE
- VIII ELONGATION FACTOR G (EF-G)
- IX ELONGATION FACTOR TU (EF-TU)
- X 30S RIBOSOMAL PROTEIN S12
- XI 50S RIBOSOMAL PROTEIN L12
- XII 50S RIBOSOMAL PROTEIN L14
- XIII VALYL tRNA SYNTHETASE (VALRS)

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XIV CELL DIVISION PROTEIN FtSH HOMOLOG

XV DnaK PROTEIN (HSP70)

XVI GTP BINDING PROTEIN LepA

XVII TRANSPORTER and

XVIII OLIGOPEPTIDE TRANSPORT ATP BINDING PROTEIN OPPF.

- 7. (Previously Presented) A method as claimed in claim 1 wherein the said method of comparing the peptide libraries as given in step (iii) of claim 1 is carried out by following the steps:
 - selecting organism names from a menu;
- iteratively comparing peptide sequences of a first organism to peptide sequences of a second organism and for matching sequences, writing sequences to a file for the first organism and to a file for the second organism.
- 8. (Currently Amended) A method as claimed in claim 1 wherein the said method of locating the common peptides in the original protein sequences as given in step (iv) of claim 1 is carried out by following the steps:
 - selecting protein sequences;
 - iteratively comparing matched peptide sequences to protein sequences;
- if the peptide is found in a protein sequence, <u>labeling</u> the peptide sequence in a file associated with the protein with: a) a protein identification number (PID), b) a location in the protein sequence, and c) a name of the organism.

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9. (Previously Presented) A method as claimed in claim 1 wherein the said method of creating a common peptide of variable length after removing the overlapping as given in step (v) of claim 1 is carried out by following the steps:

- · iteratively comparing data on matched peptide locations;
- determining overlapping matched peptides; and
- determining extended peptide sequences based on overlapping matched peptide sequences.
 - 10. (CANCELLED)
 - 11. (CANCELLED)
 - 12. (CANCELLED)